List of Subjects in 16 CFR Part 305

Advertising, Energy conservation, Household appliances, Labeling, Reporting and recordkeeping requirements.

Accordingly, 16 CFR Part 305 is amended as follows:

PART 305—[AMENDED]

1. The authority citation for Part 305 continues to read as follows:

Authority: 42 U.S.C. 6294.

2. Appendix F to Part 305 is revised to read as follows:

Appendix F to Part 305—Clothes Washers

Range Information:

''Compact'' includes all household clothes washers with a tub capacity of less than 1.6 cu. ft. or 13 gallons of water.

''Standard'' includes all household clothes washers with a tub capacity of 1.6 cu. ft. or 13 gallons of water or more.

Capacity	Range of estimated annual energy consumption (kWh/yr.)	
	Low	High
Compact:		
Top Loading	592	607
Front Loading	(*)	(*)
Standard:		
Top Loading	294	1231
Front Loading	241	318

(*) No data submitted.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 98–10391 Filed 4–17–98; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF THE TREASURY

Customs Service

19 CFR Part 142

[T.D. 98–34]

Technical Correction Regarding Time Limit for Filing Documentation After Release

AGENCY: United States Customs Service, Department of the Treasury.

ACTION: Final rule.

SUMMARY: This document makes a technical correction to the regulation regarding time limit for filing documentation after release, in accordance with Customs policy of periodically reviewing its regulations to ensure that they are current and accurate.

EFFECTIVE DATE: April 20, 1998.

FOR FURTHER INFORMATION CONTACT: Harold Singer, Chief, Regulations Branch, (202) 927–2268.

SUPPLEMENTARY INFORMATION:

Background

In accordance with Customs policy of periodically reviewing its regulations to ensure that they are current and accurate, Customs has discovered that there is a typographical error in § 142.23, Customs Regulations (19 CFR 142.23). Section 142.23 was amended by Treasury Decision 80–26, which was published in the **Federal Register** (45 FR 3901) on January 21, 1980. When that amendment was codified in Title 19 in 1980, the word ''period'' was inadvertently omitted. The error has been carried forward in each volume of Title 19 since that publication. This document corrects the error.

Inapplicability of Public Notice and Comment and Delayed Effective Date Requirements, the Regulatory Flexibility Act, and Executive Order 12866

Inasmuch as this amendment merely corrects a typographical error, pursuant to 5 U.S.C. 553(b)(B), notice and public procedure thereon are unnecessary, and pursuant to 5 U.S.C. (d)(3), a delayed effective date is not required. Since this document is not subject to the notice and public procedure requirements of 5 U.S.C. 553, it is not subject to the provisions of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*). This amendment does not meet the criteria for a ''significant regulatory action'' as defined in Executive Order 12866.

Drafting Information

The principal author of this document was Janet L. Johnson, Regulations Branch, U.S. Customs Service. However, personnel from other offices participated in its development.

List of Subjects in 19 CFR Part 142

Customs duties and inspection.

Amendment to the Regulations

Part 142, Customs Regulations (19 CFR part 142), is amended as set forth below.

PART 142—ENTRY PROCESS

1. The general authority citation for part 142 continues to read as follows:

Authority: 19 U.S.C. 66, 1448, 1484, 1624.

§ 142.23 [Amended]

2. Section 142.23 is amended by removing the words ''for quota class merchandise within the quota'' and adding in their place ''for quota class merchandise within the quota period''.

Dated: April 15, 1998.

Harold M. Singer,

Chief, Regulations Branch.

[FR Doc. 98–10316 Filed 4–17–98; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 610

[Docket No. 97N–0449]

RIN 0910–ZA08

Revisions to the General Safety Requirements for Biological Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Direct final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the biologics regulations by adding ''cellular therapy products'' to the list of products excepted from the general safety test (GST) and by adding an administrative procedure for obtaining exemptions from the GST requirements for other biological products. FDA is taking this